AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Currently Amended) ~~Pharmaceutical~~ **A dry pharmaceutical** preparation comprising a **dry** mixture of powdery teicoplanin and at least one powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin, amikacin, tobramycin, vancomycin, moxifloxacin and ciprofloxacin and an inorganic and/or organic adjuvant.

2. (Currently Amended) ~~Pharmaceutical~~ **The dry pharmaceutical** preparation pursuant to claim 1, which contains calcium carbonate, calcium sulfate dihydrate, tricalcium phosphate and/or hydroxylapatite as the inorganic adjuvant.

3. (Currently Amended) ~~Pharmaceutical~~ **The dry pharmaceutical** preparation pursuant to claim 1, which contains polyesters of at least one of lactic acid, glycolic acid, 5-hydroxy valeric acid, 6-hydroxy caproic acid and co-polymers thereof as organic adjuvants.

4. (Currently Amended) ~~Pharmaceutical~~ **The dry pharmaceutical** preparation pursuant to claim 1, which is in the form of one of tablets, molded bodies, fibers and granules.

5. (Currently Amended) ~~Pharmaceutical~~ **The dry pharmaceutical** preparation pursuant to claim 1, comprising a combination of polymerizable methacrylic acid esters and mixtures consisting of powdery teicoplanin and at least one powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin, amikacin, tobramycin, vancomycin, moxifloxacin and ciprofloxacin formed and polymerized into a molded body.

6. (Currently Amended) ~~Pharmaceutical~~ **The dry pharmaceutical** preparation pursuant to claim 1, wherein the mixture is part of a resorbable and/or of non-resorbable coating, which has been applied to non-metallic and metallic implants.

7. (Currently Amended) ~~Pharmaceutical~~ **The dry pharmaceutical** preparation pursuant to claim 1, wherein before being cured inorganic calcium phosphate bone cements and plaster mixtures are admixed to mixtures consisting of powdery teicoplanin and at least one powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin, amikacin, tobramycin, vancomycin, moxifloxacin and/or ciprofloxacin.

8. (Currently Amended) A permanent or temporary implant comprising a **dry** pharmaceutical preparation pursuant to claim 1 in the form of one of tablets, molded bodies, fibers and granules.

9. (Currently Amended) A method of treating a bacterial infection in a patient in need thereof comprising administering to said patient a pharmaceutical preparation pursuant to claim 1.

10. (New) A method of preparing the dry pharmaceutical preparation according to claim 1, said method comprising dry mixing powdery teicoplanin and at least one powdery, water soluble salt form of at least one of gentamicin, clindamycin, kanamycin, amikacin, tobramycin, vancomycin, moxifloxacin and ciprofloxacin and an inorganic and/or organic adjuvant.

CONDITIONAL PETITION FOR EXTENSION OF TIME

If entry and consideration of the amendments above requires an extension of time, Applicants respectfully request that this be considered a petition therefor. The Commissioner is authorized to charge any fee(s) due in this connection to Deposit Account No. 14-1263.

ADDITIONAL FEE

Please charge any insufficiency of fees, or credit any excess, to Deposit Account No. 14-1263.